PRISCA 5.0.2.37

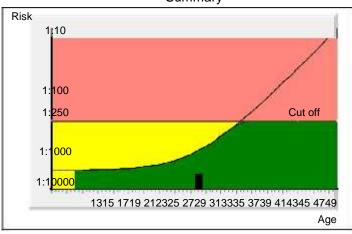
Licenced for: KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Results for:	Sample no	Date of report:
MRS. RANBIR KAUR	1710220158/CMP	11/10/17

Referring Doctors

Summary



Patient data	a
Age at delivery	32.1
WOP	19+ 5
Weight	73.6 kg
Patient ID	

For MRS. RANBIR KAUR, born on 16-01-1986, a screening test was performed on the 10-10-2017. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

 Value
 Corr. MoMs

 AFP
 57.1 ng/ml
 0.97

 HCG
 16908 mIU/ml
 1.02

 uE3
 1.4 ng/ml
 0.97

Gestation age 19+ 5
Method BPD Hadlock

The MoMs have been corrected according to:

maternal weight ethnic origin

	Risks at term	
1	Biochemical risk for Tr.21	1:2515
	Age risk:	1:725
	Neural tube defects risk	<1:10000

TRISOMY 21 SCREENING

The calculated risk for Trisomy 21 is below the cut off which represents a low risk.

After the result of the Trisomy 21 test it is expected that among 2515 women with the same data, there is one woman with a trisomy 21 pregnancy and 2514 women with not affected pregnancies.

The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.

Please note that risk calculations are statistical approaches and have no diagnostic value!

TRISOMY 18 SCREENING

The calculated risk for trisomy 18 is < 1:10000, which indicates a low risk.

NEURAL TUBE DEFECTS (NTD) SCREENING

The corrected MoM AFP (0.97) is located in the low risk area for neural tube defects.





